ARSIANOVA, A.Kh.; BELYAKOV, V.D.; BERGER, B.I.; VASIL'YEV, A.S.; GAVRILOV,
N.A.; GELMAN, L.I.; KALUGIN, V.P.; KOROSTELEV, V.Ye.; K-AMER,
I.I.; MIKHAYLOVSKIY, V.T.; E.OGOZIN, I.I.; SEREERYAKOV, L.V.

Combined vaccination with chemical and living vaccines. Voen.-med.

(MIRA 14:2)

(VACCINATION)

MIKHAYLOVSKIY, V.T., general-mayor meditsinskoy sluzhby; AGAFONOV, V.N.,

polkovnik meditsinskoy sluzhby, dotsent

Enidemiological nature of influenza and problems in its prevention.

Voen.-med. shur. no.3:77-82 Mr '60.

(INFLUENZA)

(INFLUENZA)

ROGOZIN, Isuak Iosifovich, red.; BELYAKOV, V.D., red.; KOLOSTELEV, V.Ye., red.; MINIAYLOVSKIY, V.T., red.; SOLODILOV, Ye.V.. red.; LABEZOV, G.T., red.; SHURA-BURA, B.L., red.; DAAL'-BELG, I.I., red.; LEBEDEVA, Z.V., tekhn. red.

[Military epidemiology]Voennaia e; ideniologiia. Leningrad, Yedgiz, 1962. 135 p. (MIRA 15:11)

(EPIDEMIOLOGY) (MEDICINE, MILITARY)

MIKHAYLOVSKIY, V. T., and AGAFONOV, V. I.

"Some results and problems with respect to further lowering the incidence of infectious disease among troops" - p. 3

Voyenno Meditsinskiy Zhurnal, No. 3, 1962

USSR / Form Animals. Cattle

Q-2

Abs Jour : Rof Zhur-Biol., No 6, 1958, 26131

Author

: Mikhaylovskiy V.V.

Inst

: Not given

Titlo

The Effect of the Rations containing Coriander Grist upon the Productivity of Cows, Quality of Milk and of the Butterfat (Experimental Work). (Vliyaniye ratsioner s keriandrevym shrotom na produktivnosť korov, kachestve meloka i maslyanego zhira (Eksperim, rabota))

Orig Pub: Tr. Voronozhsk. zoovot. in-ta, 1956, 14, 53-62

ibstract: The coriander grist consists of ripe coriander grains from which husk, othereal oil and fat have been eliminated.

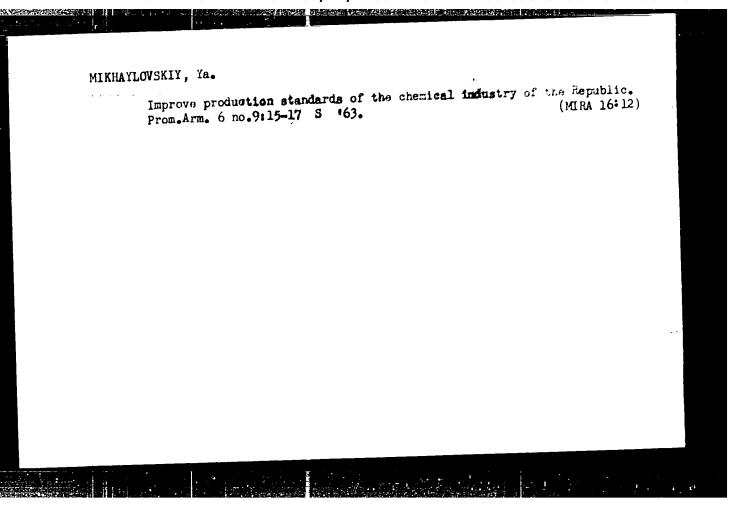
It contains 12.1 percent of hygroscopic water, 25 percent of protein, 5.6 percent of fat, 19.2 percent of cellulose, 29.2

Card 1/2

14

MIKHAYLOVSKIY, V. V., Candidate Med Sci (diss) -- "The nervous intermesentery tract of man and certain mammals (Experimental-morphological investigation)". Odessa-Lugansk, 1959. 15 pp (Odessa Med Inst im N. I. Pirogov, Lugansk Med Inst), 200 copies (KL, No 24, 1959, 151)

# Improve the supply of chemical materials to Armenian industry. Prom.Arm. 6 no.1:15-17 Ja '63. (MIRA 16:4) (Armenia-Chemical industries)



MIRGABEROV, G.G., inzh.; MIKHAYLOVSKIY, Ye.A., kand. tekhn. nauk

Review of F.B. Karpin's book "Design of weighing and proportioning equipment." Mekh. i avtom. proizv. 18 no.4:58 Ap'64.

(MIRA 17:5)

ACC NRI AP6021577 SOURCE CODE: UR/0402/66/000/003/0328/0332 (N)AUTHOR: Mikhaylovskiy Ye. M. ORG: Institute of Poliomyelitis and Virus Encephalitis Discases, Academy of Medical Sciences, SSSR, (Institut poliomiyelita i virusnykh entsefalitov AMN SSSR) TITLE: Growth of rabies virus in primary puppy kidney tissue culture SOURCE: Voprosy virusologii, no. 3, 1966, 328-332 TOPIC TACS: virology, virus, rabies wires, production method, tissue culture, serial passage, animal disease, human disease, disease vector ABSTRACT: A comparative study of two strains ("Mochalin" and "Kap") of rables virus grown in puppy kidney tissue showed that the "Mochalin" strain which had been adapted to the culture by serial passaging multiplied more rapidly than the "Kap" strain. No destruction of the cell monolayer was observed during multiplication of the virus. The "Mochalin" strain had also been adapted to growth in a hamster kidney cell culture and multiplied rapidly when transferred to the puppy kidney culture. Orig. art. has: 2 tables. [W.A. 50; CBE No. 10] SUB CODE: 06/ SUBM DATE: 15Jul65/ ORIG REF: 004/ OTH REF: 003/ UDC: 576.858.21-095.6.093.35

MIKHAYLOVSKIY, Yevgeniy Vasil'yevich, kandidat tekhnicheskikh nauk, dotsent; LiZO,G.F.. redaktor: KOGAN,F.L., tekhnicheskiy redaktor

[Theory and design of the automobile] Teoriia i raschet avtomobilia. Moskva, Nauchno-tekhn. izd-vo avtotransp. lit-ry, 1955.

246 p. (MIRA 9:2)

(Automobiles--Design and construction)

MINIMATION Transfer Vestligavich, kandidat tekhnicheskikh nauk, dotsent;

GENZINOV, V.I., redaktor; GALARTICKOVA, Ye.F., tekhnicheskiy redaktor;

[The construction of sutomobiles] Ustroistvo avtomobilia. Koskva,

Wauchno-tekhn. izd-vo avtotransp. lit-ry, 1956. 337 p. (MLRA 10:1)

(Automobiles-Design and construction)

MIKHAYLOVSKII, Tevgeniy Vasil'yevich; TSIHBALIN, Viktor Borisovich;
SHKCL'NIKOV, A.B., red.; PEVZNER, V.I., tekhn.red.

[Theory of tractors and sutomobiles] Teoriis traktors i
svtomobilia. Moskva, Gos.izd-vo sel'khos.lit-ry, 1960. 335 p.

(MIRA 13:11)

(Tractors) (Automobiles)

MIKHAYLOVSKIY, Yevgeniy Vasil'yevich; SAFONOV, Georgiy Anatol'yevich; SEREBRYAKOV, Kirill Borisovich; TARASOVA, K.A., red.; YUNISOVA, K.I., tekhn. red.

[Motor-vehicle and tractor engines] Avtotraktornye dvigateli.
Gor'kii, Gor'kovskoe knizhnoe izd-vo, 1963. 302 p.
(MIRA 17:4)

GRIGOR'YEV, E.P., inzh.; KUZNETSOV, V.Ye., inzh.; MAKSHEYEV,
V.O., inzh.; FETROVSKIY, A.S., inzh.; VEDESHKIB, V.I.,
tekhnik; KORAEEL'NIKOV, V.V., kapitan-nastavnik;
MIKHAYLOVSKIY, Ye.V., red.

[Fisheries] Promyslovoe delo. Murmansk, Murmanskoe knizhnoe
izd-vo, 1964. 463 p. (MIRA 18:5)

15(2)
AUTHORS: Gorodinskiy, G. M., Kudryashov, A. M., Mikhaylovskiy, Yu. E.

TITLE: New Models of Reflexometers (Novyye modeli refleksometrov)

PERIODICAL: Steklo i keramika, 1959, Nr 7, pp 37 - 39 (JSSR)

THE THE MENTERS OF

As may be seen from papers by G. M. Gorodinskiy, A. G. Minakov, R. I. Tsoy (see footnote) up to now the attachment reflexometer NRG-1 was used for the operational control of the working accuracy of flat polished surfaces in glass works. This device exhibits a number of shortcomings and must be operated by 2 persons. Figure 1 shows the new model of the attachment reflexometer RN1 which may be operated by one person. Its wiring diagram is represented in figure 2. For this purpose the valves 6N9S, the ferroresonance voltage stabilizer STN-35M, the germanium diodes of the type DGTs-27 and the incandescent lamp STs61 were used. For the purpose of examining the accuracy of the processing of glass which afterwards is to be polished, a test sample of a recording reflexometer RR1 was produced

type M24, and the electronic potentiometer of the type EPP-09. The reversible motor DT-75 and the electromagnetic muff

(see figures 3 and 4). Its measuring device consists of the antimony-cesium-photocell STsV-6, the microamperemeter of the

Card 1/2 EMR-500 are fitted into the carriage mechanism. The recording

Mew Models of Reflexometers

SCY/72-59-7-12/13

reflexometer was tested and mounted in the polishing line of the Gusevskiy Glass Works. The calibration of the reflexometer is carried out according to the GOST 2789-51. By means of this device only clean, degreased, and dry glass shall be examined. There are 4 figures and 2 Soviet references.

Card 2/2

L 10222-66

ACC HR. AF5028497

SOURCE\_CODE: UR/0286/65/000/020/0074/0074

AUTHOR: Mikheylovskiy, Yu. K.

26

ORG: none

TITLE: Monochromator. Class 42, No. 175680

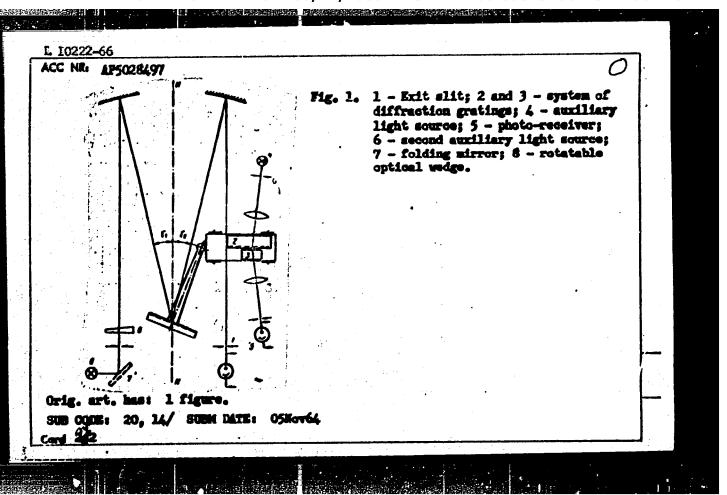
SOURCE: Byulleten' isobreteniy i tovarnykh snakov, no. 20, 1965, 74

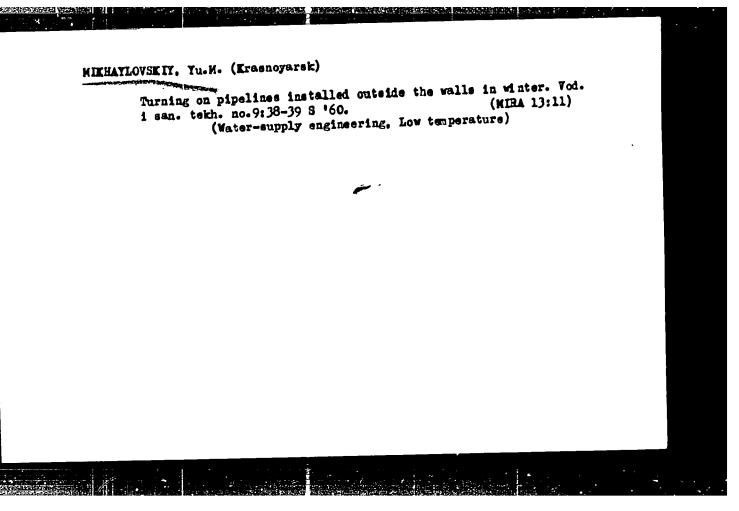
TOPIC TAGS: monochromator, optics, optical instrument, diffraction grating

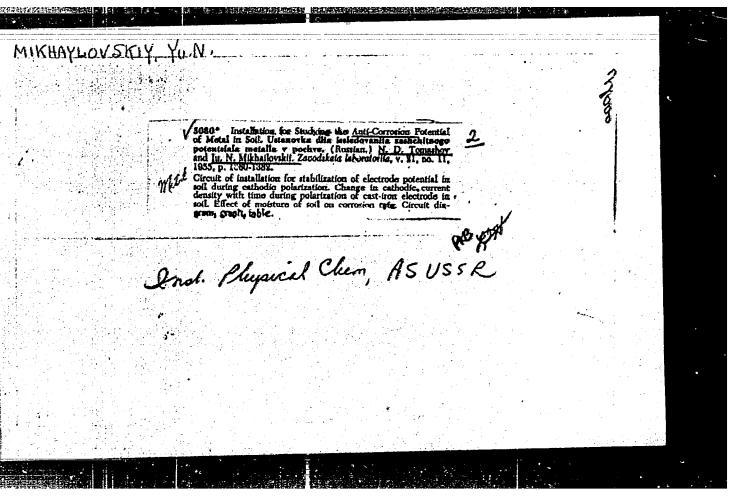
ABSTRACT: This Author Certificate describes a monochromator with a horizontal-symmetrical optical scheme, containing a diffraction grating-turning mechanism. To increase the accuracy and automation of the process of directing the spectral lines on to the exit slit, the monochromator is equipped with a system of measuring diffraction gratings which are coupled to the sinusoidal mechanism. The latter form, with the aid of an auxiliary light source, an interference pattern on a photoreceiver. To compensate for errors arising as a result of deformations in the framework of the device, the latter is equipped with a correction system (see Fig. 1). The system consists of a second auxiliary light source, a folding mirror which directs the auxiliary light flux along the direction of the main flux, and a rotatable optical wedge. In addition, one or both slits of the device are made movable.

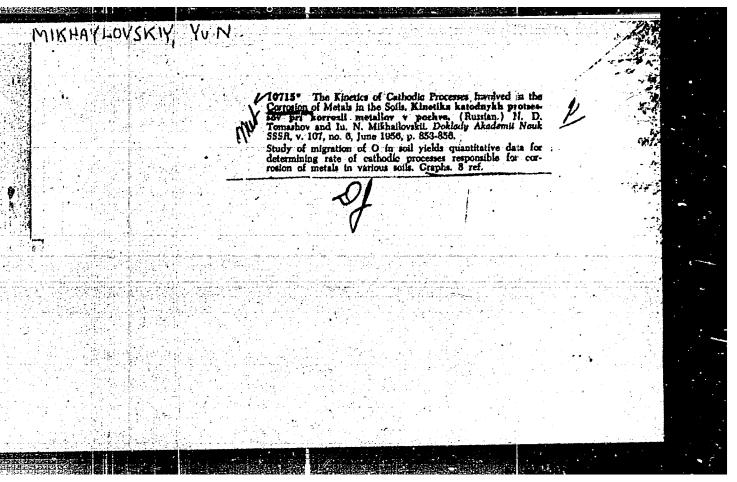
Cord 1/2

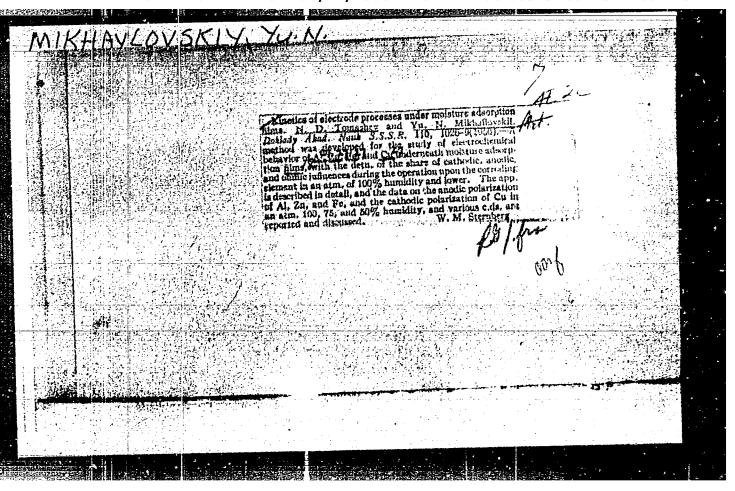
VDC: 535.853.34











USSR/Corrosion - Protection From Corrosion

J.

: Referat Zhur - Khimiya, No 4, 1957, 14090

Author

Abs Jour

Tomashov N.D., Mikhaylovskiy Yu.N.

Inst Title

: Academy of Sciences USSR : Mechanism of Electrochemical Corrosion of Metals on the

Soil

MICHAYLOV SMIY, YU. IV.

Orig Pub

: Dokl. AN SSSR, 1956, 108, No 4, 668-671

Abstract

In expanding the concepts of an electrochemical mechanism of corrosion in the soil there are pointed out the following characteristic types of macrocorrosion couples, the existence of which often determines the most pronounced destruction of metallic underground structures. I. Macrocouples, due to different permeability to oxygen, of soil of different nature (clay, dand). 2. Macrocouples associated with local heterogeneity of soil. 3. Macrocouples formed as a result of boundary effect of the diffusion of 02 in the soil at the edges of the

Card 1/2

- 2 -

APPROVED FOR RELEASE: 06/14/2000

CIA-RDP86-00513R001134110014-5"

MIKHAYLOVSKIY, Yu. N. Cand Chem Sci.,

MIKHAYLOVSKIY, Yu. H., Cand Chem Sci., -- (diss) "Study of the electrochemical processes of soil corrosion of metals." Mos, [1957]. 16 pp. (Acad Sci USSR, Inst of Phys Chemistry), 100 copies. (KL, 9-58, 113)

- 20 -

MIKHHYLEVSKID TO 11

AUTHORS:

Kikhaylovskiy, Yu.N., Tomashov, N.D.

32-12-29/71

TITLE:

The Method of the Electrochemical Investigation of Atmospheric Corrosion in the Case of a Relative Air Moisture of 100% and Less (Metod elektrokhimicheskogo isaledovaniya atmosfernoy korrozii pri otnositel'noy vlazhnosti vozdukha ot 100% i nizhe).

PERIODICAL:

Zavodskaya Laboratoriya, 1957, Vol. 23, Nr 12, pp. 1462-1466 (USSR)

ABSTRACT:

In the introduction it is said that the quantitative data on the kinetics of electrochemical processes taking place under a moisture-adsorption layer are still lacking in scientific publications. Therefore such a method is suggested. For this purpose 2 foil strips of the metals to be investigated (length 100 mm, thickness 50°), which serve as electrodes, were placed between three glass plates of equal size (and 1.5 mm thickness), and were pressed together in a plexiglass stand. The front surfaces formed by the ends of the glass- and metal strips were ground and formed the working surfaces on which the wires of the line were led to the metal plates. All this was placed into an exsicutor, and on the upper surfaces of the plexiglass stand two moist strips of filter paper were placed, to which two copper sulphate electrodes for comparison were added. In the interior of the exsicutor a certain moisture content of the air (saturated with acid

Card 1/7

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The Method of the Electrochemical Investigation of Atmospheric Corrosion in the Case of a Relative Air Moisture of 100, and Less

32-12-29/71

solution) was oreated, viz. a) moisture 100%, h) moisture 75% with NaCl content, and c) moisture 50% with Ca(NO3)2 content. The electric part of the exsicoator further consisted of a galvanometer, a battery of 150 V, a changeable resistance of 1.5-2.00 mg ohm, and one for 100,000 chm, to this cathode a voltmeter was connected, the wiring circuit of which is described separately. Electric wiring makes it possible to maintain regulated polarization; the film of moisture forming on the front surfaces of the electrodes serves as electrolyte. In this connection it is pointed out that only the polished front surfaces of glass with metal had the property of adsorbing the layer of moisture, because such a layer does not form on plexiglass. Therefore, also the other surfaces (with the exception of the front surfaces) of the glass plates were coated with paraffin, so that the aforementioned adsorption on these surfaces became impcasible. A number of experiments carried out in this connection is mentioned, the results of which are shown in form of drawings and in tables. There are 5 figures, 2 tables, and 7 Slavic references.

Card 2/3

The Method of the Electrochemical Investigation of Atmospheric Corrosion in the Case of a Relative Air Moisture of 100% and Less 32-12-29/71

ASSOCIATION: Institute for Physical Chemistry AN USSR (Institut fizicheskoy

khimii Akademii nauk SSSR).

AVAILABLE: Library of Congress

Card 3/3 1. Atmospheric corrosion 2. Moisture-Air relativity

3. Electrochemical-Determination

MIRHATLOVSKIY, Yu.B.; TOMASHOV, H.D.

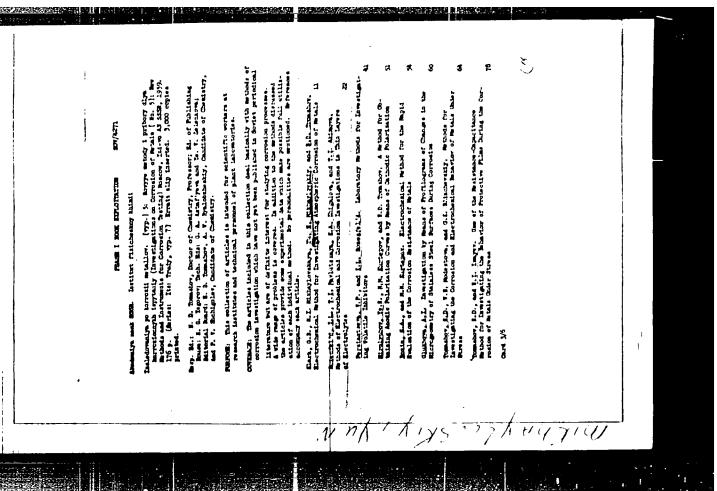
Method for field investigation of corrosive properties of soils.
Zav. lab. 23 no.4:450-454 '57. (MIRA 10:6)

1. Institut fizicheskoy khimii Akademii nauk SSSR. (Soils—Analysis) (Electrolytic corrosion)

TOWASHOV, W.D., prof., doktor khim. nauk, otvetstvennyy ied.; YEESHOV, I.K., kand. tekhn. nauk, red.; LUMEV, A.F., kand. khim. nauk, red.; NIEHAYLOVSKIY, Yu.M., kand. khim. nauk, red.; STRIZHEVSKIY, I.V., kand. tekhn. nauk, red.; SHCHIGOLEV, P.V., kand. khim. nauk, red.; BANKVITSER, A.L., red. izd-va; KASHIWA, P.S., tekhn. red.

[Theory and practice of corrosion protection for underground installations; papers of the Sixth All-Union Conference on Corrosion and Protection of Metals] Teoriia i praktika protivo-korrosionnoi zashchity podsemnykh scoruzhenii; trudy VI Vsesoiuznego zovezhchaniia po korrosii i zashchite metallov. Moskva, Izdvo Akad. nauk SSSR. 1958. 273 p. (MIRA 11:10)

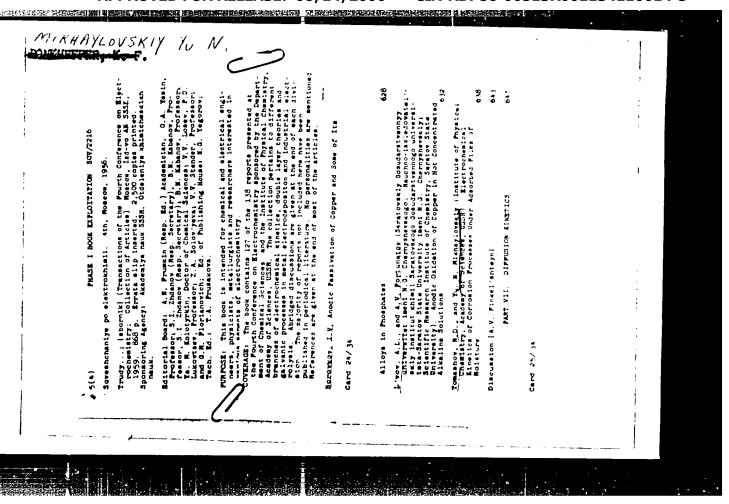
1. Vsesoyusnoye soveshchaniye po teorii i praktike protivokorroziomoy zashchity podzemykh soorusheniy. 6th, 1956. (Electrolytic corrosion)



TOMASHOV, Mikon Danilovich, Prinimali uchestiye: TYUKINA, M.N.; PALEOLOG, Ye.N.; CHERNOVA, Q.P.; MIKHAYLOVSKIY, Yh.N.; LUNEY, A.P.; TIMONOVA, M.A.; MODESTOVA, V.T.; MATYETEVA, T.V.; BYALOEREKKIY, A.V.; ZHUK, M.P.; SHRETDER, A.V.; TIYOV, V.A.; VERNETEVA, M.A.; LOKOTILOV, A.A.; EXRUKSHTIS, Q.K.; DERYAGINA, Q.Q.; FEDOTOVA, A.Z.; FOKIN, M.N.; MIROLTUBOV, Ye.N.; ISATEV, N.I.; AL'TOVSKIY, R.M.; SHCHIGOLEV, P.V., YEGOROV, N.G., red.izd-va; KUZ'MIN, I.F., tekhn.red.

[Theory of the corrosion and the protection of motals] Teorise korrozii i zashchity metallov. Moskva, Izd-vo Akad.nauk SSSR, 1959. 591 p. (MIRA 13:1)

(Corrosion and anticorrosives)



MIKHAYLOVSKIY, Yu.N.; TOMASHOY, N.D.

Using the method of polarization by intermittent currents in the investigation of corrosion processes in atmospheres with high ohmic resistance. Trudy Inst.fiz.khim. no.7:85-95 '59.

(MIRA 13:5)

(Blectrolytic corrosion) (Polarization (Electricity))

SUV/20-124-6-29/55 5(2,4) AUTILE: Penashov, E. D., Mishaylevskiy, Yu. N. Mechanism of Anodic Dissolution of Metals in Soils (Mekha-TITLE: nizm anodnogo rastvoreniya metallov v pochvakh) Doklady Akademii nauk SSSR, 1959, Vol 124, Nr 6, PELICOICAL: pp 1285 - 1288 (USUR) For the dissolution mentioned in the title as well as for ADSTLACT: electrolytes "e general equation holds: Me " mH20 + ne, in which the primary stage of the process is a migration of the metal ion into the soil electrolyte. As far as the anodic process is accompanied by a hydration of the forming metal ions the presence of a certain amount of moisture in the soil is an indispensable condition (Ref 1) Greater variations in the moisture of natural soils may have a considerable influence upon the rate of anodic metal dispolution. In this connection the authors mention the investilation results of the aforesaid process to the abe of "Armko" iron in soils of different Ca-4 1/7

Mechanism of Anodic Dissolution of Metals in Soils

SCV, 20-124-6-29/55

moisture Figure 1 gives anodic polarization curves obtained in the case of iron electrodes in sand with 1-20% moisture in which case the moisture was added in form of fall solution. From the results it can be seen that the stable potential of iren is shifted into the positive range in the case of decreasing soil moisture; the inhibitions of the anodic reaction increase. A similar dependence exists in loam oils from the results obtained it can be seen that the density of the self-dissolution currents increases with decreasing soil me, ture, i.e in connection with making less complicate the cathodic process (Ref 1) whereas, the stable potential of iron is shifted on the general curve 1 into the positive range (Fig. ). Thus, the corresion rate inreen as an included for the active (just moistened) anodic surface with operanding buil measture. If the this rate is calculated for the visible surface the former will intrease only as long as the making less complicated at the catholic process proceeds more rapidly than the shrinking of the active surface. In the case of a further decrease in moisture the entire rate of metal corrosion will decrease in convenience of the pastivation of the basic surface of

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the metal. There are f figures, 1 table and Colevier forences.

AND CHARION: Institut finicheskey Minima Akademii nauk SUSR (Institute of Physical Chemistry of the Abberry of Sciences, USR)

PRESENTED: November 6, 1978, by V. I. Spitsyn, Academician

SUBMITTED: November 7, 1980

ISAYEV, N.I.; Prinimali uchastiye: MIKHAYLOVSKIY, Yu.N.; BERUKSHTIS, G.K.

Atmospheric corrosion of steel wire reope. Trudy Inst.fiz.khim.
8:144-154 '60. (MIRA 14:4)

(Wire rope—Corrosion)

18 8300

27215 S/081/61/000/014/013/030 B103/B217

AUTHORS:

Tomashov, N. D., Mikhaylovskiy, Yu. N.

TITLE:

Electrochemical theory of underground corrosion of metals

PERIODICAL:

Referativnyy zhurnal. Khimiya, no. 14, 1961, 332, abstract 140182. (Tr. In-ta fiz. khimii. AN SSSR, 1960, vyp. 8.

190 - 216

TEXT: The authors examined data on the effect of soil structure and properties (humidity, permeability) on cathodic and anodic processes in underground corrosion. It was found that the rate of uniform total corrosion I of a metal in the soil can be calculated from the equation  $I_{\text{corr}} = KI_KI_a / (I_K + I_a), \text{ where } K \text{ is a constant, } I_K \text{ is the density of the cathodic limiting current at an iron electrode in the soil concerned (oxygen permeability of soil), } I_a \text{ is the density of the anodic current at a given potential. The rate of local corrosion ( ) can be estimated from the oxygen permeability of soil <math>I_K$  and from its resistance ( ) :  $AI_K /$  ,

Card 1/2

Electrochemical theory of ...

27215 S/081/61/000/014/013/030 B103/B217

where A is a constant. The possibility of formation of large macrofields is determined by the equation  $I = B\Delta I_K/\Delta I_f^{-1}$ , where B is a constant,  $I_K/\Delta I_f$  is the change of oxygen permeability in a soil section of length  $\Delta I$ . On the basis of these assumptions a device for estimating the corrosive activity of soils was designed and constructed. [Abstracter's note: Complete translation.]

Card 2/2

TOMASHOV, N.D.; MIKHAYLOVSKIY, Yu.N.; LEONOV, V.V.

Investigating the work of afferential aeration couples in soils.
Trudy Inst.fiz.khim. 8:217-225 '60. (MIRA 14:4)

(Soil corrosion)

TOMASHOV, N.D.; KRASNOYARSKIY, V.V.; MIKHAYLOVSKIY, Yu.N.

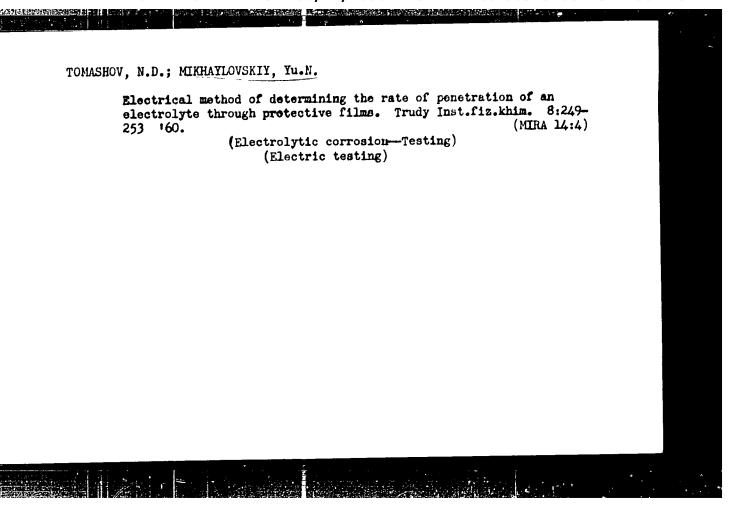
Field testing of the corrosion of steels insoils. Trudy Inst.fiz.
khin. 8:226-234 \*60. (MIRA 14:4)

(Steel--Corresion) (Soil corrosion)

TOMASHOV, N.D.; LUNEV, A.F.; MIKHAYLOVSKIY, Yu.N.; LEOKOV, V.V.

Determination of protective properties of metal coatings. Trudy Inst.fiz.khim. 8:235-248 '60. (MIRA 14:4)

(Protective coatings)



TOMASHOV, N.D.; MIKHAYLOVSKIY, Yu.N.; LOPOVOK, G.G.

Testing of insulation coatings for cracking during flaxure. Trudy
Inst.fiz.khim. 8:276-280 '60. (MIRA 14:4)

(Protective coatings—Testing)

TOMASHOV, N.D.; MIKHAYLOVSKIY, Yu.N.; LEONOV, V.V.

Kinetics of the deterioration of portective coatings on motals in electrolytes. Trudy Inst.fiz.khim. 8:291-296 '60. (NIRA 14:4)

(Protective coatings) (Electrolytic corrosion)

TOMASHOV, N.D.; MIKHAYLOVSKIY, Yu.N.; LEONOV, V.V.

Kinotics of cathodic processes in the corrosion of metals under protective coatings. Trudy Inst.fiz.khim. 8:297-304 '60.

(Protective coatings) (Electrolytic corrosion)

(Protective coatings)

29420

18 8300

S/081/61/000/017/058/166 B110/B138

AUTHORS:

Tomashov, N. D., Mikhaylovskiy, Yu. N., Leonov, V. V.

TITLE:

Study of the action of macrocorrosion pairs formed when a metal surface is partially protected by thin protective films

PERIODICAL:

Referativnyy zhurnal. Khimiya, no. 17, 1961, 293, abstract

1711218 (Tr. In-ta fiz khimii. AN SSSR, no. 8, 1960.

305 - 312)

TEXT: The authors investigated the effect of macrocorrosion pairs on protected and unprotected metals immersed in an electrolyte consisting of 0 5 N NaCl, 0.016 N  ${\rm H_2O_2}$  and 0.01 N HCl at  ${\sim}20^{\circ}{\rm C}$ . Protective materials

used were: asphalt, paraffin, wax, bakelite varnish, nitrocellulose, drying oil, whit with drying oil and minium with drying oil. The protective films (PF) were 1.0 - 6.0  $\mu$  thick. Irrespective of the type of the PF an insulated electrode in couple with an uninsulated one always acts as the cathode. The corrosion current of the pair increases in the order Cu-Al-Fe-Zn. The presence of an incomplete PF on the metal surface leads to an improvement in the static potential of the electrode Card 1/2

S/081/61/000 017/008.166

Study of the action of macrocorrosion... B110/B138

and the strong localization of corrosion in the pores and places where the protective film is broken. The resistance of the PF is not only dependent on the properties of the material of which it is made but also on the corrosion potential of the metal in the aggressive medium in question. Abstracter's note: Complete translation.

S/076/61/035/002/009/015 B124/B201

AUTHORS:

Tomashov, N. D., Mikhaylovskiy, Yu. N., and Leonov, V. V.

TITLE:

Mechanism of the electrochemical corrosion of metals under insulation coatings. I. Kinetics of the destruction of

insulation coatings on metals in electrolytes

PERIODICAL:

Zhurnal fizicheskoy khimii, v. 35, no. 2, 1961, 367-372

TEXT: A study has been made of the electrochemical behavior of metal electrodes insulated by means of thin bitumen-, paraffin-, bakelite-, and other coatings. The electrodes were Pt, Cu, and Fe wires, 0.5 to 1 mm in diameter. The film was applied by dipping the electrode into liquid insulating material and then slowly and uniformly extracting it by a warren motor. The thickness of the resulting film was determined by the extraction rate of the electrode and the viscosity of the insulating material. The capacity method was applied for measuring the film thickness; it amounted to  $1-4\,\mu$ . The kinetics of the destruction of the insulation film on metals in electrolytes was measured with the aid of the capacity and the resistance of the insulated electrode in a 0.5 N Card 1/5

Mechanism of the electrochemical ...

S/076/61/035/002/009/015 B124/B201

NaCl solution; the measuring device is diagrammatically shown in Fig. 1. After electrode 1 was insulated, it was dipped into glass cell 2 filled with the electrolyte. Capacity and resistance were measured with a cylindrical auxiliary Pt electrode 3. The measurement was made with the equal-armed bridge 5 which was fed by generator 4 of the type 3C2A (3G2A). The a-c amplitude did not exceed 20-25 mv. The bridge equilibrium was visually fixed with the oscilloscope 7 of the type 30-4 (E0-4), the amplifier 6 being connected to its input. Capacity and resistance were measured at determined time intervals after the electrode was dipped into the electrolyte. The tests were conducted at room temperature (20-22°C) within a maximum of 30 days. Fig. 2, a shows the curves of the change of capacity and resistance with time on a bitumen-film covered Pt electrode in 0.5 N NaCl. The capacity of the electrode rises and its resistance drops with time, which is correlated with the change of the film structure upon the action of the electrolyte. The authors theoretically infer from the measurement results that the deterioration of the insulation characteristics of coatings such as bitumen and the beginning of the corrosion process are chiefly connected with the penetration of the electrolyte into the micropores and defects of the film as far as near the metal sur-Card 2/5

Mechanism of the electrochemical ...

S/076/61/035/002/009/015 B124/B201

face, while the swelling of the film takes longer and is of no great importance. Fig. 2,6 shows that RC remains constant during a 30-day test of the Pt electrode under bitumen coatings. Fig. 3, a shows the dependence of the Pt electrode capacity under a bitumen coating on the initial frequency during 15 to 30 days. Fig. 3,6 shows the same in logarithmic coordinates. The curves of the change of capacity on Pe and Cu insulated with a thin bitumen film are given in Fig. 4. A decisive factor determining the initial rate of destruction of the insulation film is the electrochemical mature of the metal. G. V. Akimov and N. D. Tomashov are mentioned. There are 4 figures and 19 references: 7 Soviet-bloc and 3 non-Soviet-bloc. The two references to English Language publications read as follows: C. Corfield, Gas., 21, 11, 35, 1945; E. A. Koening, Oil. a.Gas J., 44, 20, 303, 1945.

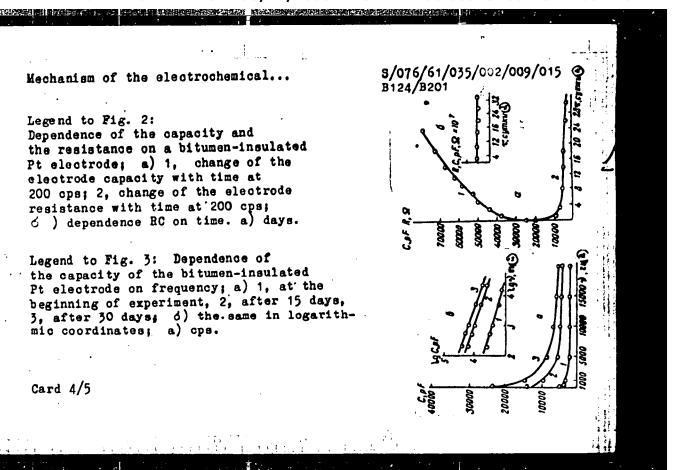
ASSOCIATION: Akademiya nauk SSSR, Institut fiziche skoy khimii

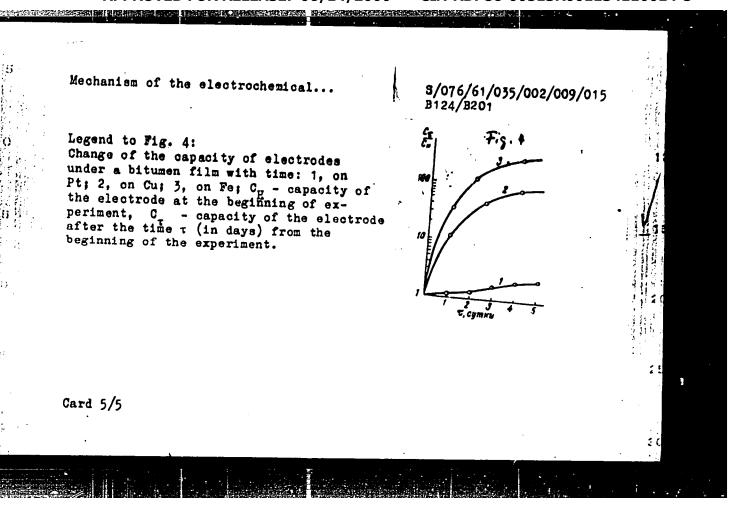
(Academy of Sciences USSR, Institute of Physical Chemistry)

· SUBMITTED:

June 2, 1959

Card 3/5





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4016, 1138, 1208

B/076/61/035/003/010/023 B121/B203

AUTHORS:

Tomashov, N. D., Mikhaylovskiy, Yu. N., and Leonov, V. V.

TITLE:

Mechanism of electrochemical corrosion of metals under insulation coatings. II. Kinetics of cathodic processes during the

corrosion of metals under insulation coatings

PERIODICAL:

Zhurnal fizicheskoy khimii, v. 35, no. 3, 1961, 588-594

TEXT: The authors studied the cathodic processes on metals (Pt, Cu, Fe) whose surface was coated with thin layers (1-6  $\mu$ ) of an insulating film (bitumen, Bakelite, and nitrocellulose lacquers) in solutions of corrosive properties. They studied the cathodic polarization and the change in capacity of insulated and noninsulated platinum electrodes in a solution of 0.1 N  $Fe^{2+}$  + 0.1 N  $Fe^{3+}$ , and observed an intense polarization at the insulated electrode, even at low cathode current density. The occurrence of electron conductivity in individual parts of the insulating film of platinum is explained with the electron conductivity in the film itself. A gradual increase of the electrochemically active metal surface occurs during the cathodic polarization, which facilitates the electrochemical process. During Card 1/2

S/076/61/035/003/010/023 B121/B203

Mechanism of ...

the electrolytic process, the layer round the electrode is in an active state promoting the cathodic process. Therefore, cathodic processes may also occur in the thinnest spots of the insulating film. The mechanism of metal corrosion under porous insulating materials was discussed. The cathodic process was assumed to take place not only on the bare metal surface but also in the finest sections of the insulating film. Macro- and microcorrosion pairs occur on the metal surface insulated with a fine porous insulating film, the free metal surface acting as anode, and the insulated part as cathode. The density of the corrosion current does not only depend on the electrochemical nature of the electrode metal but also on the electric properties of the insulating material. The authors thank A. A. Novikov for assisting in the experiments. There are 5 figures and 13 references:

12 Soviet-bloc and 1 non-Soviet-bloc. The reference to the English-language publication reads as follows: Yasushi Soto, Masuo Kamioka, Yuhei Nemoto, J. Elektrochem. Soc. Japan, 26, 1, E-26, 1958.

ASSOCIATION:

Institut fizicheskoy khimii Akademiya nauk SSSR (Institute of

Physical Chemistry of the Academy of Sciences USSR)

SUBMITTED:

July 3, 1959

Card 2/2

21997

16 8300

S/076/61/035/004/002/016 B106/B201

AUTHORS:

Tomashov, N.D., Mikhaylovskiy, Yu.N., and Leonov, V.V.

TITLE:

Mechanism of the electrochemical corrosion of metals

under insulating coatings

PERIODICAL: Zhurnal fizicheskoy khimii, v. 35, no. 4, 1961, 736 - 742

TEXT: III. Study of the mode of operation of macrocorrosion pairs forming with partial insulation of a metal surface by thin protective films

In continuation of earlier studies on the corrosion processes in metals protected by insulating coatings (Ref. 1: Zh. fiz. khimii, 3, 400, 1960; Ref. 2: Zh. fiz. khimii, 35, 367, 1961), the authors conducted a systematic investigation of the mode of operation of the macrocorrosion pair formed from an insulated and the corresponding noninsulated metal in an electrolyte solution. The specimens were prepared from wire electrodes 1 mm in diameter and 50 mm in length, made of copper, iron, aluminum, and zinc. The insulation materials used were bitumen, paraffin, bakelite lacquer, nitrocellulose, varnish, zinc white on varnish, and Fe<sub>2</sub>C<sub>3</sub> on

Card 1/6

21997

S/076/61/035/004/002/018 B106/B201

Mechanism of the electrochemical ...

varnish; the coatings were between 1 and 6  $\gamma$  thick. The method of applying the coatings has already been described in the abovementioned papers. The electrolyte solution was 0.5 N NaCl + 0.016 N H<sub>2</sub>O<sub>2</sub> + 0.01 N HCl. The

found that an insulated iron electrode always behaved as a cathode and compared to a noninsulated iron electrode, regardless of the nature of the insulation material. The maximum density of the corrosion current with iron electrodes was in most cases attained 12-15 hours after the beginning of the experiments. With equal thickness of the coating, the macrocorrosion current on an electrode with a bitumen coating was found to be 5 times as strong as on an electrode coated with varnish. The addition of pigmenting substances (excepting zinc white) to nitrocellulose and varnish effects an increase of the density of the maximum corrosion current. Similar conditions arise also in insulated zinc and aluminum electrodes. The initial capacity of insulated electrodes rises in the electrolyte solution by 2-3 orders of magnitude in the course of 48 hours, while the resistance drops to about the same extent. Such phenomena do not appear on noninsulated electrodes. The electrochemical not tree of the

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21997 \$/076/61/035/004/002/018 B106/B201

Mechanism of the electrochemical ...

electrode metal has an effect upon the corrosion current of the macrocorrosion pair. For the same insulation material (Fe<sub>2</sub>0, on varnish), the density of the maximum corrosion current grows in the succession copper < < aluminum < iron < zinc, i.e., with dropping corrosion stability of the metal in the electrolyte solution. The mechanism of corrosion under the insulating coating was also studied. A porous insulating coating on the metal surface caused the steady electrode potential to turn more positive, and corrosion to be strongly concentrated in the pores and defects of the coatings. The results of the present work show that, regardless of the nature of the insulation material, the rate of destruction of thin porous insulating coatings is in the first place dependent upon the electrochemical neture of the electrode metal and of the corrosive medium. The coating will retain its insulating properties to a degree proportional to the stability of the metal in the respective electrolyte solution. This result is of a great practical importance. N.I. Zhuravleva is thanked for her active assistance in the experimental work. The authors intend to make a special study of diffusion phenomena on insulated electrodes. There are 6 figures, 2 tables, and 4 Soviet-bloc references.

Card 3/6

oil-varnish; (8) nitrocellulose; a) hr

21997 3/076/61/035/004/002/018 B106/3201 Mechanism of the electrochemical ... Akademiya nauk SSSR Institut fizicheskoy khimii ASSOCIATION: (Academy of Sciences USSR Institute of Physical Chemistry) July 13, 1959 SUBMITTED: Fig. 2: Change of macrocorrosion FIG. 2 current with time in vapors formed from noninsulated and insulated A A/CHE iron electrodes in the solution of  $0.5 \text{ n NaCl} + 0.016 \text{ n H}_20_2 + 0.01 \text{ n}$ HCl. The insulated electrode is coated: (1) by bitumen; (2) by nitrocellulose with Cr203; (3) by bakelite lacquer; (4) by minium and oil-varnish; (5) paraffin; (6) oil-varnish; (7) zinc white and

Card 4/6

TOMASHOV, N.D., MIKHAYLOVSKIY, YU.N., LEONOV, V.V. AND HIKITENKO, YE.A.

"Electrochemical protection of buried structures from stray current corrosion by means of unilaterially polarizing anodes."

Report submitted to the Second Intl. Congress on Corresion of Metals New York City 11-15 March 1963

INSTITUTE OF PHYSICAL CHEMISTRY, MOSCOW

**治疗和抗发性的原理** 

1 35589-65 EWT(=)/EWP(b)/EWA(d)/EWP(t) JD/WB

ACCESSION NR: AT3007224

5/2951/63/000/000/0359/0367/7

AUTHOR: Mikhaylovskiy, Yu. N.; Tomashov, N. D.; Nikitenko, Ye. A.

8+1

TITLE: Unilaterally polarized metallic protectors for electrochemical prevention of corrosion gaused by stray currents in underground equipment

SOURCE: Korroziya metallov i splavov; sbornik. Hoscow, Hetallurgizdat, 1963, 359-367

TOPIC TAGS: corrosion protection, underground equipment, electrochemical process, electric drasnage

ABSTRACT: The general principles of protectors in underground corrosion prevention are studied. It is shown that a system of unilaterally polarized protectors may be used where there are relatively low stray underground currents. The corrosion prevention method which is examined is free of many of the disadvantages inherent in the electrical drainage method. The proposed system automatically draws off the current from the underground equipment in the anode zones and prevents the current in the cathode zones from reaching the equipment through the protector. The effectiveness of the method is determined by the electrochemical characteristics of the protector. Protectors made of various metals (including

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# MIKHAYLOVSKIY, Yu.N. Corrosion of iron in humid soils under the effect of an alternating current. Zhur.prikl.khim. 36 no.3:551-557 My '63. (MIRA 16:51) 1. Institut fizicheskoy khimii AN SSSR. (Iron—Corrosion) (Electric currents, Alternating)

MINHAYLOVSKIY, Yu.N.

Corrosion of lead in humid soils during polarization by alternating current. Zhur.prikl.khim. 36 no.6:1273-1278 Je '63.

(MIRA 16:8)

1. Institut fizicheskoy khimii AN SSSR.

(Lead--Corrosion) (Polarization (Electricity))

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S/076/63/037/001/012/029 B101/B186

AUTHOR:

Mikhaylovskiy, Yu. N. (Moscow)

TITLE:

Electrochemical mechanism of the corrosion of metals effected by alternating current. I. Corrosion and electrochemical

behavior of iron in electrolytes on polarization by

alternating current

PERIODICAL: Zhurnal fizicheskoy khimii, v. 37, no. 1, 1963, 132-137

TEXT: The polarization curves of Fe in 0.5 N NaCl and 0.1 N HCl+0.4 N NaCl were plotted on polarization by d-c or sinusoidal a-c of 50 cps. In neutral and acid solutions, a-c polarization was much lower than d-c polarization. It is assumed that the forward reaction Fe Fe<sup>2+</sup> + 2e proceeds easily during the anodic half-period and the reverse reaction proceeds easily during the cathodic half-period. Corrosion of Fe in neutral or alkaline solutions is low, as the iron ions formed in the anodic half-period are reprecipitated during the cathodic half-period.

From the decrease in phase shift angle from  $80^{\circ}$  at 5 ma/cm<sup>2</sup> to  $65^{\circ}$  at Card 1/3

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Electrochemical mechanism of the ...

60 ma/cm<sup>2</sup>, a current 1 of the double layer charge reversal is calculated to be 54 ma/cm<sup>2</sup> for 60 ma/cm<sup>2</sup>, and a Faraday current i<sub>r</sub> is calculated to be 25  $ma/cm^2$ . In acid media, the reverse course of the cathodic polarization curve does not coincide with the direct course. Hence, the electrode surface is assumed to change. For a square-wave a-c at 5 cps, the cathodic polarization curves in 0.5 N NaCl, 0.5 N NaCl + 0.1 N KOH, and 0.4 N NaCl + 0.1 N HCl at an anodic current density  $i_{\perp} = 12 \text{ ma/cm}^2$  were plotted to eliminate the effect of  $i_c$ . Results: at  $i_c < i_a$ , the cathodic potential is a linear function of  $\log 1_c$ . As soon as  $i_c > 1_a$ , the potential of Fe rapidly shifts into the discharge region of hydrogen ions. The effect of the HCl concentration on the dissolution rate of iron was examined in 1 N NaCl for a square-wave at 5 cps a-c, with [HCl] being changed from  $10^{-4}$  to 1 N. Up to  $10^{-3}$  N HCl, the dissolution rate increased but slightly. Between  $10^{-3}$  and  $5 \cdot 10^{-2}$  N HCl, the dissolution rate was proportional to the acid concentration. Conclusions: The dissolution rate of Fe on a-c polarization in non-passivating media depends on the rate of Card 2/3

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Electrochemical mechanism of the ...

the cathodic depolarization processes. The dissolution rate is accelerated by the presence of oxidants reducing on the iron at a potential more positive than that of the reversible iron electrode. There are 4 figures.

ASSOCIATION: Akademiya nauk SSSR, Institut fizicheskoy khimii

(Academy of Sciences USSR, Institute of Physical

Chemistry)

SUBMITTED: September 30, 1961

Card 3/3

### "APPROVED FOR RELEASE: 06/14/2000

CIA-RDP86-00513R001134110014-5

S/076/63/037/002/008/018 B101/B186

AUTHOR:

Mikhaylovskiy, Yu. N. (Moscow)

TITLE:

Electrochemical mechanism of metal corrosion influenced by alternating current. II. Dissolution of lead on

polarization with alternating current

PERIODICAL:

Zhurnal fizicheskoy khimii, v. 37, no. 2, 1963, 340-346

TEXT: Studies were made on the corrosion of lead due to polarization by a 50-cps sinusoidal or a 5-cps square-shaped a-c in 0.5 N NaCl, 1 N KNO3,

0.001 - 1 N H<sub>2</sub>SO<sub>4</sub>, 0.001 - 1 N HNC<sub>5</sub>, and 1 N CH<sub>5</sub>COOH using a lead electrode rotating at 2500 rpm. Results: In general the corrosion rate increased with the density of the a-c and depended on the nature of the medium. With 50 cps a-c, the double-layer recharging current was only 17-20% of the polarization current. In neutral media the corrosion rate of Pb was low owing to the periodical ionization and precipitation of its ions. The rate of dissolution in neutral media depended on the cathodic oxygen depolarization which proceeds in parallel with the decharging of the Pb ions:

Card 1/2

s/076/63/037/002/006/016 B101/B166

Electrochemical mechanism of ...

 $C_2 + 2H_2O \longrightarrow 40H^ (i_{0_2})$  and tended toward a constant value limited by the diffusion rate of the oxygen. In  $H_2SO_4$  the rate of corrosion was almost independent of the acid concentration within the range from 0.001 to 1 N. In oxidizing acids, the rate of corrosion was proportional to the acid concentration. For  $HNO_3$ ,  $i_{corr} = A \left[ HNO_3 \right]^{0.5}$ ; A is a constant. With frequencies of more than 5 cps, the metal ions forming in the anodic halfperiod can no longer diffuse through the diffusion layer so that the total corrosion effect is determined mainly by the kinetics of the cathodic processes. There are 5 figures.

ASSOCIATION:

Akademiya nauk SSSR, Institut fizicheskoy khimii (Academy of Sciences USSR, Institute of Physical

Chemistry)

SUBMITTED:

November 4, 1961

Card 2/2

## s/076/63/037/003/005/020 B101/B215

AUTHOR:

Mikhaylovskiy, Yu. N.

TITLE:

Electrochemical mechanism of corrosion of metals under the

action of a-c. III. Dissolution of zinc during

polarization by a-c

PERIODICAL: Zhurnal fizicheskoy khimii, v. 37, no. 3, 1963, 553-558

TEXT: The electrochemical and corrosive behavior of zinc was studied in 0.5 N NaCl or 0.1 N HCHO.4 N NaCl at room temperature under the action of 50 cps isinuscidal a-c or 5 cps restangular a-c. Results: (1) The rate of Zn dissolution by a-c in a neutral solution at a current density up to 30 - 40 ma/cm² during polarization depends on the rate of cathodic reaction of oxygen ionization; at higher current densities it depends on the discharge of H<sup>+</sup> ions. (2) In an acid solution, the rate of Zn dissolution depends mainly on the discharge of H<sup>+</sup> ions in the cathodic half-period. (3) In general the rate of metal dissolution in acid solutions is proportional to the current density of polarizing a-c. In Card 1/2

Electrochemical mechanism of ... S/076/65/037/003/005/020

Electrochemical mechanism of ... B101/B215

a weakly acid solution the dissolution rate increases with the H<sup>+</sup>
congentration. In strongly acid solutions this dependence no longer exists.

(4) The self-dissolution of Zn explains the higher rate of Zn dissolution in acid solutions as compared with the value calculated theoretically. There are 4 figures.

ASSOCIATION: Akademiya nauk SSSR Institut fizicheskoy khimii (Academy of Sciences USSR Institute of Chemical Physics)

SUBMITTED: November 21, 1961

#### "APPROVED FOR RELEASE: 06/14/2000

CIA-RDP86-00513R001134110014-5

L 9905-63 EWP(q)/EMT(m)/EDS\_AFFTC/ASD\_JD/JG

ACCRESSION HR: AP3000430 8/0076/63/037/005/1196/1200

55

AUTHOR: Mikhaylovakiy, Yu. H. 53

TITIE: Electrochemical mechanism of metal corrosion by alternating current.

4. Dissolution of aluminum and magnesium during polarization by alternating current

SOURCE: AN SSSR. Zhurnel fizicheskoy khimii, v. 37, no. 5, 1963, 1196-1200

TOPIC TAGE: aluminum, magnesium, polarization by alternating current, anode, ionization reaction

ABSTRACF: The electrochemical and corrosive behavior of aluminum and magnesium during polarization by alternating current in various electrolytic media was studied. The dissolution rate of these metals under the effect of alternating currents of slight density depends upon the resistance of the protective film in the given medium. The dissolution rate of aluminum and magnesium does not depend upon the hydrogen ion concentration when large densities of polarizing current are used. The breaking of the protective film on aluminum in an anode

Card 1/2

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ACCESSION NR: AP3000430

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half-period is determined by the duration of the anode half-period an magnitude of electrode potential. The dissolution rate of aluminum and magnesium during polarization by alternating current is determined by the rate of the anode ionization reaction. "Author sincerely thanks Professor N. D. Tomashor for his constant interest, attention and valuable advice during the writing and discussion of this article." Orig. art. has: 6 figures.

ASSOCIATION: Akademiya nauk SSSR (Academy of Sciences, SSSR) Institut Fizicheskoy Khimii (Institute of Physical Chemistry)

SUPPLYTHED: 23MOV61 DATE ACQ: 18Jun63 KNCL: 60

SUB CODE: 00 MR REF SOV: 006 OTHER: 000

Card 2/2

APPROVED FOR RELEASE: 06/14/2000

CIA-RDP86-00513R001134110014-5"

S/020/63/148/003/029/037 B101/B186

AUT HOR :

Mikhaylovskiy, Yu. N.

TITLE:

Card 1/2

Dissolution and passivation of titanium in sulfuric acid during polarization with alternating current

PERIODICAL: Akademiya nauk SSSR. Doklady, v. 148, no. 3, 1963, 617 - 619

TEXT: The processes underlying the passivation of titanium were experimentally studied. Titanium was passivated in 10 N sulfuric acid with an asymmetric square-shaped 10-cps alternating current. The cathodic current density  $i_{\rm C}$  was 20 ma/cm<sup>2</sup> and was kept constant; the anodic current density  $i_{\rm C}$  was varied between 0 and 20 ma/cm<sup>2</sup>. The dissolving rate of Ti and its passivation were determined. Results: Increasing  $i_{\rm R}$  improved the potential of Ti during the anodic half-cycle. The anodic potential was stabilized for all  $i_{\rm R}$ ; thus, the anodic reactions were not inhibited up to a potential of +1.0 v. In the cathodic half-cycle, the delay in the discharge of H ions depended on the preceding anodic potential in spite of  $i_{\rm C}$  const. A

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Dissolution and passivation of ...

positive shift of the ancdic potential by +0.2 v shifted the cathodic potential by -0.12 to -0.15 v each. At an anodic potential of +0.2 v, Ti ionization rapidly decreased owing to the formation of a passivating layer, and Ti dissolution ceased at +1.0 v. Conclusion: The passivating layer consists of oxygen adsorptively bound to the metal but again released in the cathodic half-cycle, and of more firmly bound oxides which increase the H overvoltage and are slowly and incompletely dissolved during the cathodic half-cycle. In the anodic half-cycle the defective spots first affected in the passivating layer are those on which adsorbed oxygen was reduced. The bonds in the passivating layer are rearranged with increasing occupation of the electrode by adsorbed oxygen. The proportion of oxide compounds in the passivating layer depends on the anodic potential. Thus titanium is passivated by an adsorptive layer in which oxygen is bound in two ways. There are 3 figures.

ASSOCIATION: Institut fizicheskoy khimii Akademii nauk SSSR (Institute of .... Physical Chemistry of the Academy of Sciences USSR)

PRESENTED: May 12, 1962, by A. N. Frumkin, Academician

SUBMITTED: October 12, 1962

Card 2/2

ENF(q)/ENT(m)/EDS AFFTC/ASD JD/WB L 12/18-63 S/0020/63/150/00L/0852/ ACCESSION NR: AP3001112 AUTHOR: Tomashov, N. D.; Strukov, N. M.; Mikhaylovskiy, Tu. N. TITIE: The effect of alternating current frequency on the speed of corrosion of titanium in sulfuric acid AN SSSR. Doklady, v. 150, no. 4, 1965, 852-855 TOPIC TAUS: electrochemical properties of titanium, corrosion properties of SOURCE: titanium, polarization of titanium, titanium ABSTRACT: The investigation of the electrochemical and corrosion behavior of titanium during its polarization with a sinusoidal variable current frequency showed that, with an increase of frequency of the polarizing current, the polarizing capability of the electrode in the anodic and in the cathodic half-period is decreased. This is additionally connected with the presence of current capacity and the explanation of easier electrochemical reactions on the surface of the electrode in the anodic and cathodic half-periods. The corrosion of titanium at lower frequencies and low densities of the polarizing current is greater than the corrosion at higher frequencies. However, a reverse effect is observed with very high densities of the polarizing current. The

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ACCESSION NR: AF3001412

examination of experimenting material shows that the variable current intensifies the process of anodic dissolution of titanium as a result of the destruction of the passive state of the electrode in the cathodic half-period. It appears that at high frequencies of the variable current in the anodic half-period when the surface of metal contains mainly the adsorbed oxygen, the activation of metal in the cathodic half-period is much greater, and thus, the dissolution of titanium is greater. At low frequencies the metal surface is covered mainly with the chemically bound oxygen in the form of oxide layers which is not completely reduced in the cathodic half-period and therefore the active surface of the metal is smaller and the speed of dissolution of the metal is smaller. Orig. art. has: 5 graphs.

ASSOCIATION: Institut fizicheskoy khimii Akademii nauk SSSR (Institute of Physical Chemistry of the Academy of Sciences SSSR)

ENGL: 00 DATE ACQ: 01Ju163 SUBMITTED: 29Jan63

OTHER: 000 NO REF SOV: 003 SUB CODE: 00

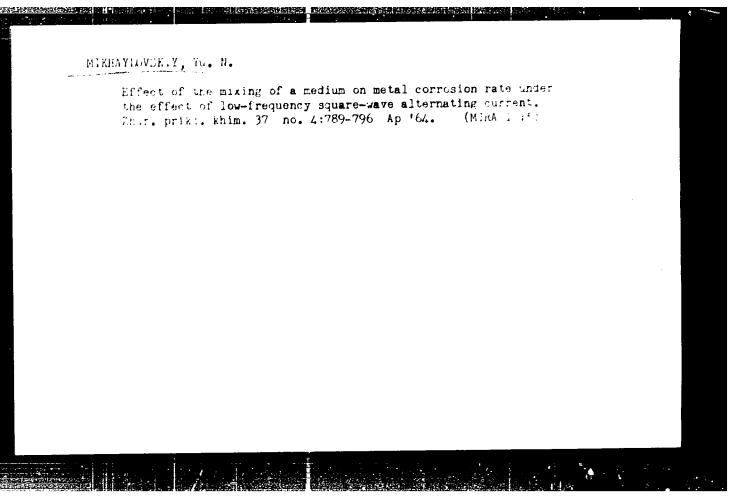
2/2 Card

> CIA-RDP86-00513R001134110014-5" APPROVED FOR RELEASE: 06/14/2000

MIKHAYLOVSKIY, Yu.N.

Corrosion of fully polarized two-electrode systems under the effect of alternating current. Zhur.prikl.khim. 37 no.1:118-126 Ja '64. (MIRA 17:2)

1. Institut fizichesko/ khimii AN SSSR.



1 17526-65 ENT(in)/ENA(d)/ENP(t)/ENP(b) ASD(f)-2/ASD(m)-3/AFMDC MJH/JD/HM/WB

ACCESSION NR: AP5000512 \$/0080/64/037/011/2528/2530

AUTHOR: Mikhaylovskiy, Yu. N., Lopovok, G. G., Tomashov, N. D.

TITLE: Corrosion of IKhl8N9T stainless steel under the influence of an

alternating current

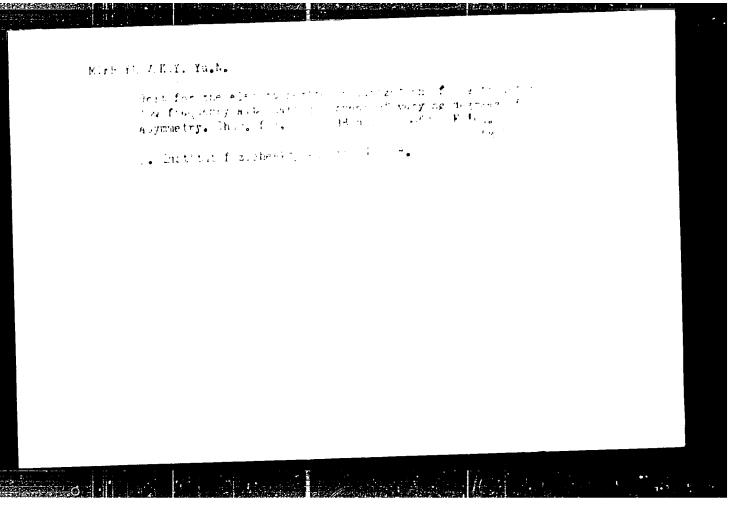
SOURCE: Zhurnal prikladnoy khimii, v. 37, no. 11, 1964, 2528-2530

TOPIC TAGS: steel corrosion, polarization corrosion, stainless steel, steel rassivation, steel 1Kh18N8T

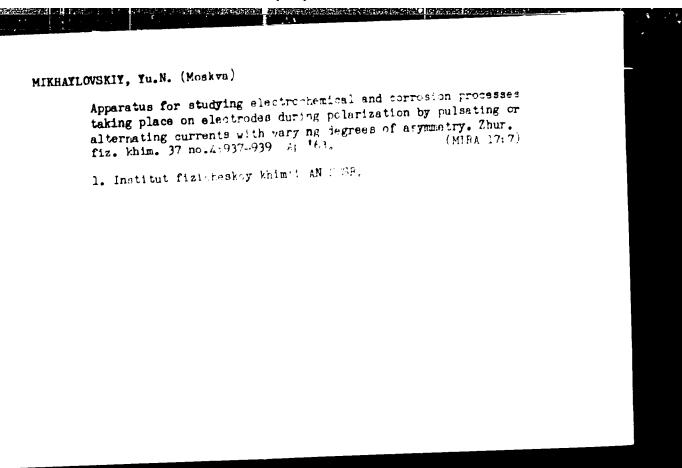
ABSTRACT: Studies on the electrochemical behavior of 1Kh18N9T steel indicated that the corrosion rate during polarization with a 50-cycle current in neutral and acid solutions is determined basically by the rate of the cathodic reactions of oxygen ionization and hydrogen ion discharge. With a given alternating current density, polarization increases with the hydrogen ion concentration in the solution. No passivation occurs, even in 10 N H2SO4, during alternating current polarization with densities up to 100 ma/cm2 because the passivating layer does not have enough time to form during the anodic semicycle. Orig. art. has 3 graphs.

Card 1/2

ASSOCIATION: Institut fizicheskoy khimii AN SSSR (Institute of Physical Chemistry, AN SSSR)  SUBMITTED: 13Dec62 ENCL: 00 SUB CODE: MM  NO REF SOV: 005 OTHER: 004	L 17526-65 ACCESSION NR: AI5000512					
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ACCESSION NR: AP4034586

5/0076/64/038/004/0995/0998

AUTHOR: Mikhaylovskiy, Yu. N.

TITLE: Corrosion of metals by the action of alternating current and agitated

media.

SOURCE: Zhurnal fizicheskoy khimii, v. 38, no. 4, 1964, 995-998

TOPIC TAGS: corrosion, alternating current, diffusion, diffusion layer boundary, polarization, agitation, agitation rate, Faraday's law

ABSTRACT: The test method described previously by the author (Zh. fiz. khimii, 37, 946, 1963) was used in this study of the effect of the rate of rotating the disc electrode on the corrosive current density formed by polarization by alternating current frequencies of 0.04-50 cycles/second. The corrosion studies were run on iron in a solution of 0.5N NaCl + 0.01N HCl + 0.0N Fe+3 and on lead in 1N KNO3. Equilibrium was established in the systems in 1 to 0.1 seconds as the electrode rotation was increased from ; w >0 revolutions/second. At frequencies below 5 cycles the corrosive effect on the metal was determined primarily by the diffusion of the metal ions in the electrolyte past the boundary of the diffusion layer

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ACCESSION NR: AP4034586

during the anodic half cycle of the current. Then the rate of corrosion approached that calculated theoretically according to the Faraday value. When the frequency of the polarizing current was more than 5 cycles, an insignificant amount of the metal ions succeeded in diffusing (during the anodic half cycle) past the limits of the diffusion layer. This was confirmed by the insignificant corrosion of metals upon polarization with alternating current of more than 5 cycles in media containing no oxidants. The increased corrosion on the metals with increased agitation at frequencies > 5 cycles/sec. was attributed to the alleviated transport of the depolarizer which is reduced on the electrode in the cathodic half cycle. Decreasing the agitation of the media shifted the frequency limit to lower values. Orig. art. has: 4 figures.

ASSOCIATION: Akademiya nauk SSSR (Academy of Sciences SSSR) Institut fizicheskoy khimii (Institute of Physical Chemistry)

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NO REP SOV: 005

OTHER: 000

Card 2/2

8/0076/64/038/004/1001/1003

ACCESSION NR: AP4034587

AUTHOR: Mikhaylovskiy, Yu. N.

TITIE: Rectification of alternate current and metal corresion in multiple electrode systems

SOURCE: Zhurnal fizicheskoy khimii, v. 38, no. 4, 1964, 1001-1003

TOPIC TAGS: multiple electrode system, current rectification, metal corrosion, oxide layer, aerating property, corrosion rate, polarization, anode dissolution

ABSTRACT: Rectification of the alternate current and increase of the corrosion process in such systems are not always determined by the aerating properties of the oxide layers; the non-linear character of the polarization curves and their asymmetry with respect to the corrosion potential, both in anode and cathodecontrolled systems, are the main determinants. The effect of the appearance of a constant component and increase of corrosion are particularly manifest upon polarization by alternate current in corrosion systems with cathode control. In systems with enode control, Fe/Al, or Fe/Zn electrodes are quite typical for showing current rectification and increase of the corresion rate upon polarization by

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ACCESSION MR: AP4034587

alternate current. Such effect is also seen upon polarization of 2 electrodes of the same metal by direct or alternate current even in neutral media. The shift of cathode potential in the range of the hydrogen discharge reaction disrupts the symmetry of the system. Thus polarization of electrochemical systems by direct and alternate current may lead to increase of the direct current component and accelerate the process of snode, dissolution. Orig. art. has: 4 figures.

ASSOCIATION: Akedemii neuk SSSR, Institut fizicheskoy khimii (Acedemy of Sciences, SSSR, Institute of Physical Chemistry)

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SUB CODE: CC, HOK

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OTHER: OOT

| Card 2/2

ACCESSION NR: AP4041758

8/0076/64/038/006/1612/1615

AUTHOR: Mikhaylovskiy, Yu. N.

TITIE: Corrosion of titanium in sulfuric and hydrochloric acid solutions with alternating current polarization.

SOURCE: Zhurnal fizicheskoy khimii, v. 38, no. 6, 1964, 1612-1615

TOPIC TAGS: titanium, corrosion, passivation, surface property, electrochemistry, polarization

ABSTRACT: The article considers the effect of the concentration of sulfuric and hydrochloric acid on the electrochemical and corrosion behavior of titanium (VT-1) upon polarization with different frequency square wave current. The determination of the rate of corrosion of titanium showed that as long as the electrode potential remains below 0.2 v during the anodic period, the corrosion effect is proportional to the duration of the anodic half cycle. The maximum corrosion current was observed with titanium at 10 cps, and it was equal to 5 ma/cm<sup>2</sup>. It was concluded that the intense dissolution of titanium in acid solutions upon a.c. polarization results from the fact that during the cathode half cycle the passivated state of

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ACCESSION NR: AP4041758

the electrode is disrupted. In the following anodic half cycle the active metal is dissolved and again passivated. When the frequency is high the stronger oxide type films do not dissolve and thus "freezing" of these oxide residues of the passivated film takes place. The absorbed oxygen is electrochemically reduced during the cathodic half cycle. Thus, by the end of the cathodic half cycle only those passivated areas remain on the electrode in which there is a predominantly oxide type bonding. During the anodic half cycle the dissolution of metal takes place initially at the active points which occurred as a result of the reduction of the adsorbed oxygen. Orig. art. has: 4 figures.

ASSOCIATION: Institut fizicheskoy khimii Akademii nauk SSSR (Institute of Physical Chemistry Academy of Sciences SSSR)

SUBMITTED: 22Mar63

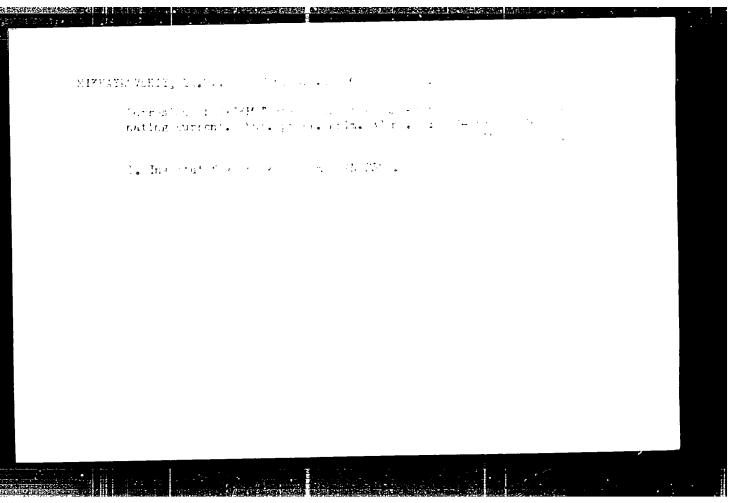
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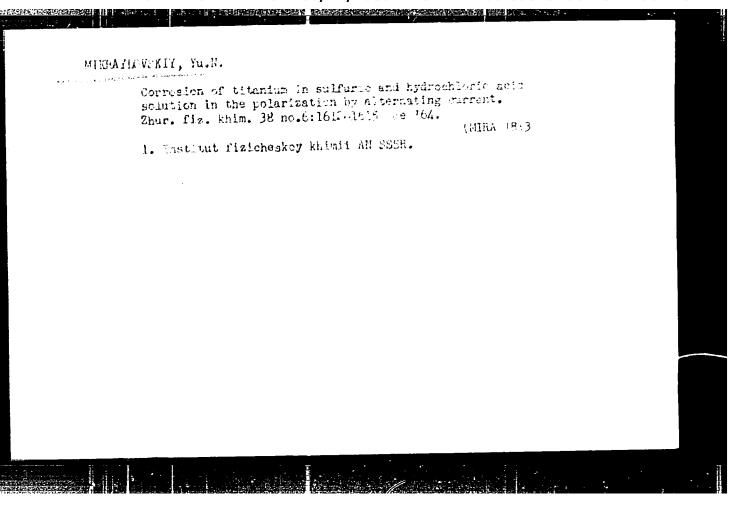
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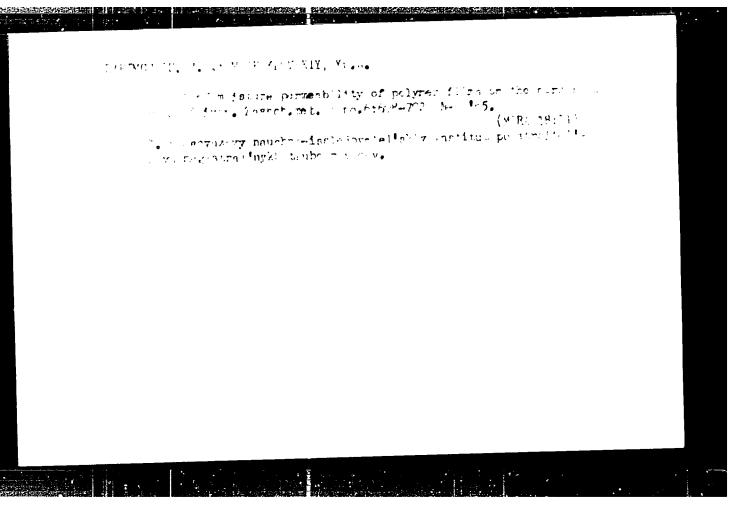




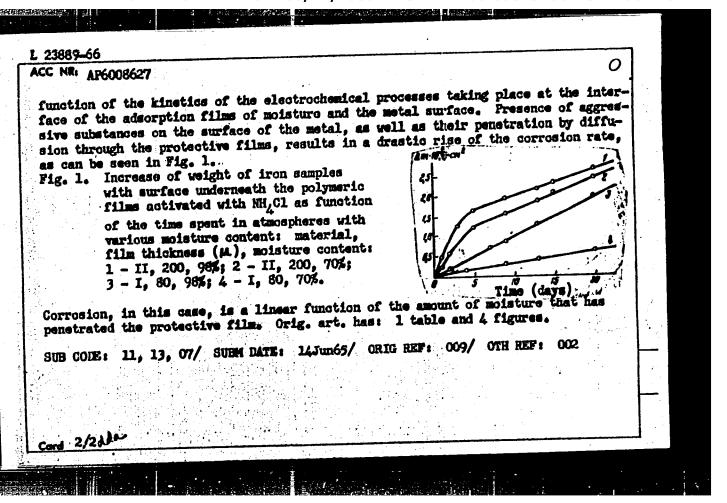
MIRHATLOVSEIT, Yu.N.; LEONOV, V.V.; TOMASHOV, N.D.

Measuring the resistance of insulating protective costings immersed in an electrolyte. Zashch. met. 1 no.51577-582 S-0 '(65. (MIRA 18:9))

1. Institut fizicheskoy khimii AN SSSR.



IJP(c) JD/NB/RM SOURCE CODE: UR/0365/65/001/006/0698/0702 AP6008627 AUTHORS: Serafimovich, V. B.; Hikhaylovskiy, Yu. N. ORG: All-Union Scientific Research Institute for Construction of Hains (Vsesoyuznyy nauchno-issledovatel'skly institut po stroitel'stvu angistral'nykh truboprovodov) TITLE: Effect of moisture permeability of polymeric films upon the iron corresion rate SOURCE: Zashchita metallov, v. 1, no. 6, 1965, 698-702 TOPIC TAGS: Acorrosion rate, corrosion protection, protective coating, polyvinyl. chloride, polyethylene plastic / V-118 polyvinyl chloride ABSTRACT: The rate of corrosion of Armco iron was investigated as a function of moisture permeability of polyethylene (I) and of polyvinyl chloride V-118 (II) covering films. The study was of considerable interest because the polymeric films are widely used as protective coverings for underground and naval metallic equipment. Moisture permeability of the films and iron corrosion (measured by increase of sample weight) were determined at 200 in an atmosphere of constant moisture content (40, 70, and 98%). It was established that the corrosion process was independent of the moisture permeability of the covering when experiments were con-2 ducted in an atmosphere of uncontaminated moisture. The process was entirely a TDC: 620.198 Cord 1/2



	L 46992-66 EMP(j)/EMT(m)/T/EMP(V) IJP(c) RM/W/ ACC NR. AP6022869 (N) SOURCE CODE: UR/0303/66/000/002/003C/0034
į	AUTHOR: Naumova, S. F.; Mikhaylovskiy, Yu. N.; Zubov, P. I.
	ORG: none
	TITLE: Effect of the vapor and gas permeability of polymer films on their properties
	SOURCE: Lakokrasochnyye materialy i ikh primenenlye, no. 2, 1966, 30-34
	TOPIC TAGS: protective coating, polymer film, hydrogen chloride, metal exidation, polyethylene, teflon, polyvinyl chloride, magnesium, ADHESIVE. BENDING
	ABSTRACT: The effect of the permeability of loose polymer film coatings on the oxidation rate of a metal in a moist atmosphere in the absence of an adhesive bond between the film and the metal was studied. The polymer films were PE-500 high-pressure polyethylene (70 \mu thick), polytetrafluoroethylene (teflon) (55 \mu), and V-118 polyvinylene (50 \mu).
	chloride (180 µ). A new method of measuring slow oxidation rates of metals was about
	of a thin metal film (~10" cm) under the polymer film. In order to increase the solu-
	sitivity of the method, the metal employed was magnesium, because of its high reactivity. It is shown that in a pure moist atmosphere the oxidation rate of the metal is
	manetically independent of the nature of the polymer film (in the case of a nonadnering)
	film). This is because the rate-determining step in the exidation is the inhibition of the anodic process of metal ionization (hydration), not the diffusion of moisture
	Card 1/2 UDC: 667.613.4